Day: Tuesday Date: 7/18/2006

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Inventor Information for 10/272929

Inventor Name	City	State/Country				
BEAUREGARD, MARC	TROIS-RIVIERES	CANADA				
GAGNON, MYLENE-CLAUDE	TROIS-RIVIERES	CANADA				
DOUCET, ALAIN	L'ANCIENNE-LORETTE	CANADA				
WILLIAMS, MARTIN	CHARNY	CANADA				
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Application Number Information

Application Number: 10/272929

Assignments

Filing or 371(c) Date: 10/18/2002

Effective Date: 10/18/2002

Application Received: 10/18/2002

Pat. Num./Pub. Num: /20030118573

Issue Date: 00/00/0000

Date of Abandonment: 00/00/0000

Attorney Docket Number: 15493-1US

PM/MG/al

Status: 41 /NON FINAL ACTION MAILED

Confirmation Number: 2858

Examiner Number: 77509 / WOITACH, JOSEPH

Group Art Unit: 1632

IFW IMAGE

Class/Subclass: 514/044.000

Lost Case: NO

Interference Number:

Unmatched Petition: NO

L&R Code: Secrecy Code:1

Third Level Review: NO

Secrecy Order: NO

Mail Non Final

Desc.

Status Date: 05/18/2006

Waiting for Response

Oral Hearing: NO Title of Invention: MB-1 ANALOGS AND USES THEREOF

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Inventor Name Search Result

Your Search was:

Last Name = BEAUREGARD

First Name = MARC

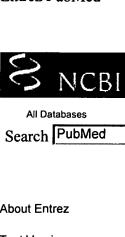
Application#	Patent#	Status	Date Filed	Title	Inventor Name
09037948	Not Issued	161	03/10/1998	GENETICALLY ENGINEERED RUMEN BACTERIAL STRAINS	BEAUREGARD, MARC
09810520	6958992	150	03/16/2001	REGISTERING AN IP PHONE WITH AN IP PHONE SWITCH	BEAUREGARD, MARC
10272929	Not Issued	41	10/18/2002	MB-1 analogs and uses thereof	BEAUREGARD, MARC
10625882	Not Issued	71	07/24/2003	MB-1 analogs and uses thereof	BEAUREGARD, MARC
<u>10776180</u>	Not Issued	71	02/12/2004	Method of mutagenic chain reaction	BEAUREGARD, MARC
11110069	Not Issued	30	04/20/2005	Registering an IP phone with an IP phone switch	BEAUREGARD, MARC
11394281	Not Issued	20	03/31/2006	System and method for scanning communications according to a policy	BEAUREGARD, MARC
60329759	Not Issued	159	10/18/2001	MB-1 analogs and uses thereof	BEAUREGARD, MARC
<u>60446518</u>	Not Issued	159	02/12/2003	Method of mutagenic chain reaction	BEAUREGARD, MARC
10306634	Not Issued	61	11/27/2002	Expandable skin for safety restraint system	BEAUREGARD, MARC P.
60278909	Not Issued	159	03/26/2001	Vehicle subfloor incorporating a ventilation system	BEAUREGARD, MARC P.

Inventor Search Completed: No Records to Display.

Last Name First Name Search Another: Inventor BEAUREGARD

MARC Search

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ClinicalTrials.gov PubMed Central	□4:	Xue HH, Kovan Brady JN, Leon		ise-Masison CA,	Berg M, Rado	ovich MF,	Related A	Articles, Link	(S
		T lymphocyte Proc Natl Acad	es. Sci U S A	lates IL-7 rece A. 2002 Oct 15;99 led - indexed for I	- 9(21):13759-6	_			l
	□ 5:	Moore SA, King Baker HM, Bake		, Loomes KM, Ho	ernell O, Blac	kberg L,	Related A	Articles, Link	(S
		reveals bile sa loop and prov J Mol Biol. 200	alt-inde rides in: Sep 21;	cated recombing pendent conformations into hep 312(3):511-23. ed - indexed for l	rmational fl arin binding	exibility			
	□6:	Campagna S, Co	sette P,	Molle G, Gaillard	IJL.		Related A	Articles, Link	(S
		Evidence for		ane affinity of	the C-termi	nal doma	ain of bov	ine milk	

Biochim Biophys Acta. 2001 Aug 6;1513(2):217-22.

	PMID: 11470093 [PubMed - indexed for MEDLINE]	
□7:	Dopfer D, Nederbragt H, Almeida RA, Gaastra W.	Related Articles, Links
	Studies about the mechanism of internalization by mamicells of Escherichia coli isolated from persistent bovine Vet Microbiol. 2001 Jun 6;80(3):285-96. PMID: 11337144 [PubMed - indexed for MEDLINE]	
□8:	Gagnon MC, Williams M, Doucet A, Beauregard M.	Related Articles, Links
	Replacement of tyr62 by trp in the designer protein milk in significant improvement of conformational stability. FEBS Lett. 2000 Nov 3;484(2):144-8. PMID: 11068049 [PubMed - indexed for MEDLINE]	bundle-1 results
□9:	Alexander J, del Guercio MF, Maewal A, Qiao L, Fikes J, Chesnut RW, Paulson J, Bundle DR, DeFrees S, Sette A.	Related Articles, Links
	Linear PADRE T helper epitope and carbohydrate B cel conjugates induce specific high titer IgG antibody responsible of the property of the second specific high titer IgG antibody responsible of the property	
□ 10	Grundy J, Morrison JJ, MacCallum JD, Wirtanen L, Beauregard M.	Related Articles, Links
	Crystallization and stabilization of MB-1, a de novo de optimized feeding technology. J Biotechnol. 1998 Jul 30;63(1):9-15. PMID: 9764479 [PubMed - indexed for MEDLINE]	signed protein for
□11	: Grundy JE, Wirtanen LY, Beauregard M.	Related Articles, Links
	Addition of a poly-(6X) His tag to Milk Bundle-1 and pimmobilized metal-affinity chromatography. Protein Expr Purif. 1998 Jun;13(1):61-6. PMID: 9631516 [PubMed - indexed for MEDLINE]	purification using
□ 12	Delneri MT, Carbonare SB, Silva ML, Palmeira P, Carneiro- Sampaio MM.	Related Articles, Links
	Inhibition of enteropathogenic Escherichia coli adhesic by colostrum and milk from mothers delivering low-bin neonates.	
	Eur J Pediatr. 1997 Jun;156(6):493-8. PMID: 9208250 [PubMed - indexed for MEDLINE]	
□ 13	: Damsky CH, Sheffield JB, Tuszynski GP, Warren L.	Related Articles, Links
	Is there a role for actin in virus budding? J Cell Biol. 1977 Nov;75(2 Pt 1):593-605. PMID: 233748 [PubMed - indexed for MEDLINE]	
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[0118] Design Strategy

[0119] The putative modifications to MB-1 structure are illustrated in FIG. 7.

The design strategy used here focused on two aspects: 1--the restrictive effect

of a covalent bond between remote residues on the protein as a whole; and

2--the precise location of Cys which permits disulfide bridge formation. By

choosing positions as far apart as possible, one can reduce the entropy gain

upon unfolding for most of the protein. Thus, insertion of a bridge between

helices I and IV would enclose a larger part of the polypolypeptide than a

bridge involving other helices. Another consideration for using helix I is

that this MB-1 segment of sequence is sensitive to proteolytic degradation.

The restriction of helix I by Cys insertion could help prevent such a phenomenon.

[0120] The position of Cys in helices I and IV must allow sulfhydryl groups to

be properly aligned in order to minimise strain induced by bridge formation.

On the basis of geometric models built for similar proteins, it appeared that

position "d" of the heptad pattern used for MB-1 design would offer the best

geometry for bridge formation. Therefore, L13 and M87 residues were selected

for mutation to Cys. FIG. 7 depicts the expected location of the bridge in the

mutant (hereafter referred to as MB-1LH, assuming it folds as per design).

Note that for proper alignment of position "d" in helices I and IV, a left-hand

connectivity of the helices had to be assumed (i.e. the bundles are positioned

such that when helix I is at the fore front, with its N-terminus pointing down,

then helix II is placed to the left of helix I). A second scenario was

considered, in which a right-hand connectivity could be specified. Examination

of the second model in FIG. 7 suggests mutations at positions "a" in helices I

and IV, since positions "d" would be too far apart. By choosing M10

and L91

residues for mutation to Cys, we attempted to generate a mutant (named MB-1RH)

that would resemble MB-1LH as much as possible, except for reversing its $\frac{1}{2}$

connectivity.

[0121] Disulfide bridges was also inserted into MB-1Trp. This protein is a

derivative of MB-1 where Tyr62 was replaced by Trp. Position 62 in MB-1 was

chosen for the emplacement of a spectroscopic probe at the moment of initial

design. As shown on the model in FIG. 1, position 62 is part of the hydrophobic core, and a niche made of 5 Ala was built around it in order to

accommodate a larger side chain in this region of the core. The replacement of

Tyr by Trp was thought to improve on stability, and indeed, characterisation of

MB-1Trp confirmed the strategy. MB-1Trp has a melting temperature of 55.degree. C. and is more resistant to protease action than MB-1. Here we are

going to use MB-1Trp because of the increase in bulk offered by Trp in the

core, in a way to compensate for the loss of volume consequent to the mutations

used for bridge insertion.

EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	2	"20040198681".pn.	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:05
L2	2	"20040198681".pn. and MB	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:14
L3	4	"20040198681".pn. or "20030118573".pn.	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:26
L4	58237	MB- or (milk adj1 bundle)	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:27
L5	58237	(MB- or mb) or (milk adj1_bundle)	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:27
L6	410	(MB-1) or (milk adj1 bundle)	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:27
L7	4	(milk adj1 bundle)	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:28
L8	409	mb-1	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:28
L9	32	mb-1 and milk	US-PGPUB; USPAT; EPO; DERWENT	OR	OFF	2006/07/18 07:28

Woitach, Joseph

From:

Hensle, Kristine (ASRC)

Sent:

Monday, July 17, 2006 3:38 PM

To:

Woitach, Joseph

Subject:

The results have posted for SN 10/625882 seq ids 1 and an oligo search of 6, Search Acc.#

195497.

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Woitach, Joseph

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Hello,

For application: 10/625,882,

can you please do a search of SEQ ID NO: 1 (protein),

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Joe

Joseph Woitach

(571) 272-0739

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